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PATENT 671302-2007

HOV 1 0 2004 E

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

applicant(s)

: Sugamura et al.

Serial No.

10/810,527

For

NON-HUMAN MAMMAL MODEL OF AUTOIMMUNE

DISEASE HAVING OX40L GENE TRANSFERRED

THEREINTO

Filed

March 24, 2004

Examiner

TBA

Art Unit

1614

745 Fifth Avenue New York, NY 10151

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I hereby certify that this paper or fee is being deposited with the United States Postal Service "Express Mail Post Office to Addressee" Service under 37 CFR 1.10 on the date indicated above and is addressed to: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

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INFORMATION DISCLOSURE STATEMENT

Commissioner for Patents P.O. Box 1450

Alexandria, VA 22313-1450

Dear Sir:

Applicants respectfully direct the Examiner's attention to the documents listed on the accompanying PTO-1449 form, and which are enclosed herewith.

The Information Disclosure Statement submitted herewith is being filed before the mailing date of a first Office Action on the merits. Accordingly, it is believed that no fee is required. However, the Commissioner is authorized to charge any additional fee, or to credit any overpayment in fees, to Deposit Account No. 50-0320.

The filing of this Information Disclosure Statement is not an admission that the documents identified herein constitute prior art to the present application.

Consideration of this Information Disclosure Statement is respectfully requested.

Applicants respectfully request that a copy of Form PTO-1449 be initialed by the Examiner and returned to the undersigned.

Respectfully submitted,

FROMMER LAWRENCE & HAUG LLP

Attorneys for Applicants

By:

Thomas J. Kowals

Reg. No. 32,147

Angela M. Collison

Reg. No. 51,107

Tel. No. (212) 588-0800

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	AA	Brocker et al. CD4 T cell traffic control: in vivo evidence that ligation of OX40 on CD4 T cells by OX40-ligand expressed on dendritic cells in B follicles (1999) Eur. J. Immunol.29: 1610-1616						
	AB	Ndhlovu et al. Critical involvement of OX40 ligand signals in the T cell priming events during experimental autoimmune encephalomyelitis (2001) J. Immunol.167: 2991-2999						
	AC	Higgins. Regulation of T cell activation in vitro and in vivo by targeting the OX40-OX40 ligand interaction: amelioration of ongoing inflammatory bowel disease with and OX40-IgG fusion protein but not with an OX40 ligand-IgG fusion protein (1999) J. Immunol.162: 486-493						
EXAMINER				DATE CONSIDERED		-		
		rence considered, whether or not and not considered. Include cop		with MPEP 609. Draw line through mmunication to applicant.				